



# PATENT SPECIFICATION

Convention Date (United States of America): May 4, 1942.

**567,682**

Application Date (In United Kingdom): May 31, 1943.

No. 8701/43.

Complete Specification Accepted: Feb. 27, 1945.

(Under Section 6 (1) (a) of the Patents &c. (Emergency) Act, 1939, the proviso to Section 91 (4) of the Patents and Designs Acts, 1907 to 1942, became operative on Feb. 14, 1945.)

## COMPLETE SPECIFICATION

### Process for Preparing Carotenoid Concentrates from Palm Oil

We, THE PROCTOR & GAMBLE COMPANY, a corporation organized under the laws of the State of Ohio, United States of America, located in the Gwynne Building, Sixth and Main Streets, Cincinnati, Ohio, United States of America, assignees of EDDY W. ECKEY, a citizen of the United States of America, residing at 713 Reilly Road, Wyoming, in the County of Hamilton, Ohio, United States of America, do hereby declare the nature of this invention and in what manner the same is to be performed, to be particularly described and ascertained in and by the following statement:

This invention relates to the preparation of a carotenoid concentrate from palm oil, the primary object being to provide an economical process for recovering such materials in concentrated form.

A further object is to separate the carotenoid pigment substantially completely from the fatty constituent of palm oil, thereby providing a light colored fatty constituent which can be converted easily and readily into a white soap.

Numerous attempts have been made to recover carotenoid pigments from oils and fats but the processes heretofore proposed are costly and usually involve the use of complicated and unusual apparatus requiring technically trained men for operation.

We have developed a method whereby the carotenoids contained in palm oil can be concentrated economically and without the necessity of complicated or unusual apparatus, and with the added advantage that the fatty constituent of the palm oil is separated in a form easily and readily convertible into a white soap. The palm oil employed in the practice of our invention is, of course, preferably one which has not been subjected to any previous processing step which would destroy carotenoid pigment, such as a bleaching operation or high temperature.

Briefly, our method involves the treatment of palm oil, consisting mainly of relatively non-volatile fatty acid esters of glycerin, with a low molecular weight aliphatic monohydric alcohol in the presence of an alkaline alcoholysis catalyst to form volatile esters of the fatty

acid constituents and the subsequent separation of alcohol esters from the relatively non-volatile carotenoid pigments, by distillation, the conditions of time and temperature being regulated throughout the process, to avoid appreciable destruction of valuable carotenoid constituents, as follows:

Destruction of carotenoid pigment is not a function of temperature alone but of time and temperature and therefore it is impossible to give a fixed temperature limit below which no pigment is destroyed and above which substantially all is destroyed. However, in general, temperatures below 130° C. are safe even for extended periods of time such as 15 to 20 hours, temperatures in the neighbourhood of 140° C. are not detrimental under normal conditions of operation where the total time of processing is under 10 hours, and temperatures as high as 150° C. may be sustained for short periods of time such as two hours or so without serious decomposition of the carotenoid pigment.

The process will be more clearly understood from the following detailed description.

The first step of the process involves alcoholysis of a palm oil, which normally contains an appreciable proportion of carotenoid pigments. The alcohol employed for this reaction is an aliphatic monohydric alcohol, such as methyl or ethyl alcohol, having less than five carbon atoms, which will convert the palm oil glycerides into esters which can be volatilized at a temperature below that at which the carotenoid pigments will be destroyed as described above. In order to facilitate the alcoholysis reaction and thereby effect substantially complete conversion from glycerin ester to a lower alcohol ester in a relatively short time without the use of temperatures harmful to the carotenoid pigment, an alkaline alcoholysis catalyst such as sodium hydroxide, potassium hydroxide, or a preformed alkali metal methoxide or ethoxide is employed. Known acid alcoholysis catalysts such as sulfuric or hydrochloric acids are harmful to the

carotenoid pigment and are to be avoided.

Whether or not the palm oil is alkali refined to remove free fatty acids prior to alcoholysis is optional. If the oil has

5 been subjected to a refining process such as alkali refining, washing with alcohol or other process wherein the free fatty acid content is reduced to a minimum while retaining a substantial amount of 10 carotenoid pigment, then alcohol and catalyst may be added directly and the mixture heated and boiled under a reflux condenser until the desired ester interchange has been completed.

15 However, if the palm oil contains an appreciable amount of free fatty acid as in the case of crude palm oil and if for some reason it is impracticable to subject the oil to a separate processing step wherein

20 the free fatty acids are removed, then it is advisable to use in alcoholysis sufficient alkaline catalyst not only to serve as catalyst but also to neutralize the free fatty acids. Other methods of rendering

25 the free fatty acids inactive towards the catalyst such as by esterification with alcohol under suitable conditions prior to the addition of the catalyst may be practiced without departing from the spirit of our invention provided of course 30 the said methods do not destroy the carotenoid pigment in the palm oil.

The alcoholized reaction mixture comprises the acyl esters of the monohydric 35 alcohol, excess alcohol, carotenoid pigment and catalyst in solution, and liberated glycerin. Under some conditions of operation, such as in the use of

40 only a moderate excess of methyl alcohol in alcoholysis, the liberated glycerin will exist for the most part as a separate phase. Under other conditions of operation, for example those involving the use of propyl or isopropyl alcohol, and a large excess of

45 which will act as a mutual solvent, all constituents of the alcoholysis reaction mixture, including the glycerin, may be mutually soluble and exist as a single phase. In either case the reaction mixture

50 may be subjected to distillation directly in order to effect concentration of the carotenoid pigment, or it may be subjected to treatment whereby the glycerin is separated prior to distillation.

55 The latter treatment is preferred, although it is not essential in accomplishing the objects of the invention.

When it is desired to separate the glycerin prior to distillation, and in the 60 event the reaction mixture exists as a single phase with glycerin in solution, removal of this substance may be effected by water washing. If, however, glycerin exists as a separate phase, most of it may

65 be removed by decantation and any

residual glycerin, together with other water soluble substances, may thereafter be separated by washing with water.

In connection with the water washing procedure, it should be noted that the alkaline catalyst may interfere with successful operation. For example,

70 sodium methoxide will decompose in the presence of water to form methyl alcohol and sodium hydroxide. The latter compound will of course react with the palm oil fatty acid esters to form soap which

75 may develop emulsion difficulties interfering with clean separation of the aqueous phase. Accordingly, to prevent this

80 action we find it preferable to render the catalyst inactive and this is most easily accomplished by the addition of an equimolar quantity of an acid reacting

85 compound such as phosphoric acid. An alternative way of inactivating the catalyst and of simultaneously removing glycerin is to wash the mixture with aqueous alcohol. Soaps resulting from

90 the decomposition of the catalyst are thus removed and do not interfere with subsequent water washing.

If there is no objection to the presence of liberated glycerin during concentration of the carotenoid pigment by distillation of the more volatile substances, then separation of same either by decantation or by water washing may be omitted.

95 However, before such a mixture is treated to effect such concentration, it is preferable to neutralize or otherwise render inactive the alcoholysis catalyst contained therein so as to prevent reversal of the

alcoholysis reaction and the reformation 100 of the glycerides during the distillation step.

After the alcoholysis reaction mixture has been prepared for concentration either by removal of glycerin and inactivation 110 of catalyst or by inactivation of catalyst alone, it is then subjected to distillation

conditions as more fully hereinafter explained, whereby the acyl esters and other materials volatilizable under the

distillation conditions, including glycerin 115 if present, are removed and subsequently fractionated if desired and the non-volatile carotenoid pigment is recovered

in concentrated form in the distillation residue. The distilled monohydric alcohol 120 esters are almost water-white and may be employed directly in the manufacture of

white soap, for example, the alcohol being recovered for reuse if desired.

By observing the temperature restrictions hereinbefore referred to, substantially all of the original carotenoid pigment

is recovered and practically none is lost by destruction due to excessively high temperatures or prolonged heating.

70

75

80

85

90

95

100

105

110

115

120

125

130

In order that the distillation step may be conducted at temperatures within the limits set above, and in order that the monohydric alcohol ester of the fatty acid may be distilled substantially completely under these limiting temperature conditions, vacuum distillation is necessary and the use of steam or other inert gas to aid in the distillation may or may 10 not be employed as desired.

The choice of suitable vacuum conditions and the selection of suitable equipment is well within the scope of those skilled in the art. Elaborate or unusual 15 apparatus is distinctly not necessary. Any conventional type of still which is adequate for conducting simple vacuum distillations in the pressure range of 0.02 to 0.5 mm. of mercury will be suitable.

20 The following example is illustrative of the manner in which our invention may be practiced. All parts are by weight.

#### EXAMPLE

25 To 500 parts of anhydrous methanol is added 5.0 parts of sodium hydroxide pellets. After the sodium hydroxide is dissolved 1500 parts alkali-refined palm oil containing an appreciable amount of carotenoid pigment is added and the mixture heated under a reflux condenser for one hour at 70° C.

The mixture is allowed to cool and settle. The lower layer, consisting predominantly of the liberated glycerin, is decanted and the upper layer of methyl esters of the palm oil fatty acids containing the carotenoid pigment in solution is washed first with a 50:50 mixture of 30 alcohol and water and then with water alone. The washed material is then dried by warming under vacuum with a stream of nitrogen bubbling through the liquid.

40 The washed and dried solution of 45 methyl esters is then conveyed to a simple distilling vessel provided with a vapor outlet sufficiently large to avoid material back pressure in the vessel and with some loosely coiled aluminium wire in the 50 vapor space to provide some rectification of the distillate.

The distillate receiver of the distilling apparatus is connected to a conventional oil sealed rotary vacuum pump and the 55 pressure on the system is reduced to about 0.04 mm. mercury pressure. The temperature of the contents of the vessel is gradually raised by means of a suitable oil bath and distillation of most of the volatile esters takes place below 135° C. In order to remove higher boiling esters, the bath temperature is thereafter raised to 140° C. for about one-half hour, the whole period of distillation taking about 60 five hours.

Several concentrates have been prepared in accordance with our invention and we have found that it is possible to concentrate the desired carotenoid pigment in a residue amounting to two to two-and-a-half per cent of the total fat used without measurable loss of the carotenoid pigment. Furthermore, examination of the concentrates prepared in accordance with our invention has shown that no alteration of the nature of the pigment is effected when the concentrates are formed in accordance with the instant process.

70 The carotenoid concentrates have pro-vitamin A activity and are suitable for use in the preparation of pro-Vitamin A pills and prepared stock feeds. In addition, these concentrates may be employed in the manufacture of margarine as a source of coloring matter which, in addition, supplies the product with pro-Vitamin A activity.

75 In all cases the distillate which consists of the alcohol esters of the palm oil fatty acids is very light in color and if desired it may be converted into white soap in accordance with known processes without difficulty. By the employment of a small rectifying column in the distillation step, it is possible to obtain an almost water-white fatty acid product at a cost substantially less than that incurred by the distillation of the fatty acids per se, this lower cost being made possible by the 80 recovery of the carotenoid pigment which possesses an appreciable value.

85 We are aware of specification No. 531,226, which describes a process for producing concentrates of naturally occurring Vitamin E and antioxidant from a glyceride-containing vegetable oil, for example wheat germ oil, by completely substituting the glycerol portion of the glycerides in the oil with methyl-, 90 ethyl- or propyl-alcohol whereby esters of low boiling point are produced which may be distilled off under vacuum. There is no specific mention of palm oil or of the recovery of pro-Vitamin A therefrom. 95 Moreover, in the said process, as more particularly described, an acid catalyst such as hydrochloric acid is used which, as already indicated, would destroy the 100 pro-Vitamin A in the alcoholysis of palm 105 oil.

Having now particularly described and ascertained the nature of our said invention and in what manner the same is to be performed, we declare that what we 110 claim is:-

1. The process of preparing a carotenoid pigment concentrate from, palm oil, which comprises alcoholizing palm oil with a low molecular weight aliphatic 115

70

75

80

85

90

95

100

105

110

115

120

125

130

monohydric alcohol in the presence of an alkaline alcoholysis catalyst to convert free and combined fatty acids of the palm oil into volatile alkyl esters, distilling off the said alkyl esters and concentrating the carotenoid pigment in the residue of the distillation, the conditions of time and temperature being regulated throughout the process, as hereinbefore described, 10 to avoid appreciable destruction of carotenoid pigment.

2. Process as claimed in Claim 1 in which the temperature of alcoholysis and of distillation does not exceed 150° C.

15 3. Process as claimed in Claim 1 in which the temperature of alcoholysis and of distillation is not substantially higher than 140° C.

4. Process as claimed in Claim 1, 2 or 20 3 in which the aliphatic monohydric alcohol has less than five carbon atoms.

5. Process as claimed in Claim 1, 2, 3 or 4 in which unbleached palm oil is employed.

25 6. Process as claimed in any of the preceding claims in which the catalyst is rendered inactive before concentrating the carotenoid pigment by distillation of volatile constituents.

7. Process as claimed in any of the preceding claims in which liberated glycerin is removed before the distillation.

8. Process as claimed in any of the preceding claims in which the esters are washed to remove residual impurities before distillation.

9. Process as claimed in any of the preceding claims in which the alcohol is methyl alcohol.

10. Process as claimed in any of the preceding claims in which the reaction mixture, before distillation, is washed first with aqueous alcohol and then with water alone.

11. Process of preparing a carotenoid pigment concentrate substantially as described in the Example.

Dated this 31st day of May, 1943.

CARPMELS & RANSFORD,

Agents for the Applicants,

24, Southampton Buildings,

London, W.C.2.

London: Printed for H. M. Stationery Office, by Charles & Read Ltd. - 1951.

Published at The Patent Office, 25, Southampton Buildings, London, W.C.2,  
from which copies, price 2s Od. per copy, by post 2s. 1d., may be obtained.